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DETAILED ACTION

The amendment filed on July 02, 2009 has been noted, Claim 3 has been canceled. New claim 12 has been added. Claims 1-2, 4-12 are pending.

Flection/Restrictions

- In the response, Applicants insist that claims 9-10 depend on the elected claim 1, they should be rejoined with elected group I of claims.
- Upon further searching and considering the pending claims, claims 9-10 are rejoined with the elected group I.
- Claims 1-4 and 6-11 in the scope of monoclonal antibody with accession No. FERM BP-10144 are considered.
- Claim 5 is still withdrawn from consideration. Applicants are reminded that the sequence cited in claim 5 is not a further limitation of claim 1, it will not be rejoined.

Priority

 The acknowledgement is made to the English-translated foreign priority submitted by Applicants.

Claim Rejections - 35 USC § 112 (withdrawn)

The rejection of claims 4 and 7 under 35 U.S.C. 112, first paragraph has been removed in view of Applicants' assurance statement provided in the response.

Claim Rejections - 35 USC § 102

 The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- For the following rejections, Applicants are reminded: Applicant cannot rely upon the foreign priority papers to overcome this rejection because a translation of said papers has not been made of record in accordance with 37 CFR 1.55. See MPEP § 201.15.

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- 8. (Withdrawn) The rejection of claims 1-2, 6, 8, 11 under 35 U.S.C. 102(a) as anticipated by Berry et al. (J. Virol. Method July 2004, Vol. 120, pp. 87-96) has been removed in view of the foreign priority document submitted by Applicants in the response.
- 9. (Withdrawn) Claims 1-3, 6, 8, 11 are rejected under 35 U.S.C. 102(b) as anticipated by Che et al. (B) (J. Clin. Microbial, July 2004, Vol. 42, No. 6, pp. 2629-2635) under 35 U.S.C. 102(a) in light of the teaching by Marr et al. (Science, May 2003, Vol. 300, pp. 1399-1404) has been removed in view of the foreign priority document submitted by Applicants in the response.
- 10. Claims 1-3, 6, 8, 11 are rejected under 35 U.S.C. 102(b) as anticipated by Che et al. (A) (J. First Mil. Med. University 2003 Jul;23(7):640-642), which is substantiated in light of the disclosures by Rota et al. (Science 2003, Vol. 300, No. 5624, pp. 1394-1399) and US patent No. 7,547,512B2 and WO 2004085650A1 both to University of Hong Kong.
- 11. In the response, Applicants submit two major arguments in that 1). Che (A) fails to describe the SEQ ID NO: 1 and 2). The N protein used by Che (A) is not from university of Hong Kong, and 3). The virus disclosed by Marra et al. belongs to the original case cluster from Toronto, Canada. Therefore, the N protein described by Che (A) is not necessarily the same as the sequence described in the instant case.
- 12. Applicants' argument has been respectfully considered. However, although the SARS virus disclosed by Marra et al. is not used by University of Hong Kong, the argument is still not found persuasive to overcome the rejection with reasons set forth below:
- A). Che (A) does describe that the N protein used for raising the monoclonal antibody is provided by University of Hong Kong;
- 14. B). The N protein used by Che (A) is the one with 100% identical to SEQ ID NO: 1 cited in claims although they do not use the virus disclosed by Marra et al.
- 15. There are several strains of SARS viruses in the art prior to the current Application was filed. According to the instant Application, The nucleoprotein (N protein) used by Applicants to generate the monoclonal antibody is from the Urbani strain as Applicants described in the instant Application (Rota et al. Science 2003, Vol. 300, Vol. 5624), pp. 1394-1399). The N protein used by Che (A) is provided by the University of Hong Kong (See page 641, section 1.21, lines 2-3 in

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first column). Although the complete viral genomes disclosed by University of Hong Kong and the Urbani strain are not identical in full length, the two N proteins are identical as evidenced by Rota et al. and in light of the disclosures by University of Hong Kong in US patent 7,547,512B1 or WO 2004856590A1. Therefore, it is concluded that the claimed SEQ ID NO: 1 is the one having 100% identical to N protein used by Che (A).

16. In particularly, N protein is encoded by the nucleic acid sequence starting from the position 28,120 bp and ending at 29,385, but there is no difference between each other in these N protein coding regions (please see the supporting on line material by Rota et al. for Table S1 and Table S3). The two strains comprises an identical N protein in light of the disclosure by University of Hong Kong in US patent No. 7,547,512B2 (SEQ ID NOs: 2471 and 231) or WO 2004085650A1 (SEQ ID NOs: 2471, 2472) according to the sequences alignments analyses.

```
SEO ID NO 2471
   LENGTH: 1269
   TYPE: DNA
   ORGANISM: Human severe acute respiratory syndrome virus
US-7547512
 Query Match
                      100.0%; Score 1269; DB 8; Length 1269;
 Best Local Similarity 100.0%;
 Matches 1269; Conservative
                          0: Mismatches 0: Indels
                                                    0: Gaps
          1 ATGTCTGATAATGGACCCCAATCAAACCAACGTAGTGCCCCCCGCATTACATTTGGTGGA 60
Qv
Db
         1 ATGTCTGATAATGGACCCCAATCAAACCAACGTAGTGCCCCCCGCATTACATTTGGTGGA 60
         61 CCCACAGATTCAACTGACAATAACCAGAATGGAGGACGCAATGGGGCAAGGCCAAAACAG
120
Db
         61 CCCACAGATTCAACTGACAATAACCAGAATGGAGGACGCAATGGGGCAAGGCCAAAACAG
120
Οv
        180
Db
        180
        181 GGCAAGGAGGAACTTAGATTCCCTCGAGGCCAGGGCGTTCCAATCAACACCAATAGTGGT
240
        181 GGCAAGGAGGAACTTAGATTCCCTCGAGGCCAGGGCGTTCCAATCAACACCAATAGTGGT
240
Qv
        241 CCAGATGACCAAATTGGCTACTACCGAAGAGCTACCCGACGAGTTCGTGGTGGTGACGGC
```

0b 300	241	CCAGATGACCAAATTGGCTACTACCGAAGAGCTACCCGACGAGTTCGTGGTGGTGACGGC
Эў Эб0	301	AAAATGAAAGAGCTCAGCCCCAGATGGTACTTCTATTACCTAGGAACTGGCCCAGAAGCT
0b 360	301	AAAATGAAAGAGCTCAGCCCCAGATGGTACTTCTATTACCTAGGAACTGGCCCAGAAGCT
Qy 120	361	${\tt TCACTTCCCTACGGCGCTAACAAAGAAGGCATCGTATGGGTTGCAACTGAGGGAGCCTTG}$
0b 120	361	TCACTTCCCTACGGCGCTAACAAAGAAGGCATCGTATGGGTTGCAACTGAGGGAGCCTTG
2y 180	421	${\tt AATACACCCAAAGACCACATTGGCACCCGCAATCCTAATAACAATGCTGCCACCGTGCTA}$
0b 180	421	AATACACCCAAAGACCACATTGGCACCCGCAATCCTAATAACAATGCTGCCACCGTGCTA
2y 540	481	${\tt CAACTTCCTCAAGGAACAACATTGCCAAAAGGCTTCTACGCAGAGGGAAGCAGAGGCGGC}$
0b 540	481	CAACTTCCTCAAGGAACAACATTGCCAAAAGGCTTCTACGCAGAGGGAAGCAGAGGCGGC
2y 500	541	${\tt AGTCAAGCCTCTTCTCGCTCCTCATCACGTAGTCGCGGTAATTCAAGAAATTCAACTCCT}$
00 00	541	AGTCAAGCCTCTTCTCGCTCCTCATCACGTAGTCGCGGTAATTCAAGAAATTCAACTCCT
2y 560	601	$\tt GGCAGCAGTAGGGGAAATTCTCCTGCTCGAATGGCTAGCGGAGGTGGTGAAACTGCCCTC$
0b 560	601	GGCAGCAGTAGGGGAAATTCTCCTGCTCGAATGGCTAGCGGAGGTGGTGAAACTGCCCTC
Ωy 720	661	${\tt GCGCTATTGCTGCTAGACAGATTGAACCAGCTTGAGAGCAAAGTTTCTGGTAAAGGCCAA}$
0b 720	661	GCGCTATTGCTGCTAGACAGATTGAACCAGCTTGAGAGCAAAGTTTCTGGTAAAGGCCAA
2y 780	721	${\tt CAACAACAAGGCCAAACTGTCACTAAGAAATCTGCTGCTGAGGCATCTAAAAAGCCTCGC}$
0b 780	721	CAACAACAAGGCCAAACTGTCACTAAGAAATCTGCTGCTGAGGGCATCTAAAAAGCCTCGC
)y	781	${\tt CAAAAACGTACTGCCACAAAACAGTACAACGTCACTCAAGCATTTGGGAGACGTGGTCCA}$

Db 840	781	CAAAAACGTACTGCCACAAAACAGTACAACGTCACTCAAGCATTTGGGAGACGTGGTCCA				
Qy 900	841	GAACAAACCCAAGGAAATTTCGGGGACCAAGACCTAATCAGACAAGGAACTGATTACAAA				
Db 900	841	GAACAAACCCAAGGAAATTTCGGGGACCAAGACCTAATCAGACAAGGAACTGATTACAAA				
Qу 960	901	${\tt CATTGGCCGCAAATTGCACAATTTGCTCCAAGTGCCTCTGCATTCTTTGGAATGTCACGC}$				
Db 960	901	CATTGGCCGCAATTTGCACAATTTGCTCCAAGTGCCTCTGCATTCTTTGGAATGTCACGC				
Qy 1020	961	${\tt ATTGGCATGGAAGTCACACCTTCGGGAACATGGCTGACTTATCATGGAGCCATTAAATTG}$				
Db 1020	961	ATTGGCATGGAAGTCACACCTTCGGGAACATGGCTGACTTATCATGGAGCCATTAAATTG				
Qy 1080	1021	GATGACAAAGATCCACAATTCAAAGACAACGTCATACTGCTGAACAAGCACATTGACGCA				
Db 1080	1021	GATGACAAAGATCCACAATTCAAAGACAACGTCATACTGCTGAACAAGCACATTGACGCA				
Qy 1140	1081	TACAAAACATTCCCACCAACAGAGCCTAAAAAAGGACAAAAAGAAAAAGACTGATGAAGCT				
Db 1140	1081	TACAAAACATTCCCACCAACAGAGCCTAAAAAGGCAAAAAGAAAAGACATGATGAAGCT				
Qy 1200	1141	${\tt CAGCCTTTGCCGCAGAGACAAAAGAAGCAGCCCACTGTGACTCTTCTTCCTGCGGCTGAC}$				
Db 1200	1141	CAGCCTTTGCCGCAGAGACAAAAGAAGCAGCCCACTGTGACTCTTCTTCCTGCGGCTGAC				
Qy 1260	1201	$\tt ATGGATGATTTCTCCAGACAACTTCAAAATTCCATGAGTGGAGCTTCTGCTGATTCAACT$				
Db 1260	1201	ATGGATGATTCTCCAGACAACTTCAAAATTCCATGAGTGGAGCTTCTGCTGATTCAACT				
Qу	1261	CAGGCATAA 1269				
Db	1261	CAGGCATAA 1269				
W0200485650Al, SEQ ID NO: 2472 Query Match 100.0%; Score 2227; DB 1; Length 422; Best Local Similarity 100.0%; Pred. No. 2.8e-189; Matches 422; Conservative 0; Mismatches 0; Indels 0; Gaps 0;						

Qу	1	${\tt MSDNGPQSNQRSAPRITFGGPTDSTDNNQNGGRNGARPKQRRPQGLPNNTASWFTALTQH}$	60
Db	1		60
Qy 120	61	${\tt GKEELRFPRGQGVPINTNSGPDDQIGYYRRATRRVRGGDGKMKELSPRWYFYYLGTGPEA}$	
Db 120	61	GKEELFFRGQGVPINTNSGPDDQIGYYRRATRRVRGGDGKMKELSPRWYFYYLGTGPEA	
Qy 180	121	${\tt SLPYGANKEGIVWVATEGALNTPKDHIGTRNPNNNAATVLQLPQGTTLPKGFYAEGSRGG}$	
Db 180	121	SLPYGANKEGIVWVATEGALNTPKDHIGTRNPNNNAATVLQLPQGTTLPKGFYAEGSRGG	
Qy 240	181	${\tt SQASSRSSRSRGNSRNSTPGSSRGNSPARMASGGGETALALLLLDRLNQLESKVSGKGQ}$	
Db 240	181	SQASSRSSSRRGNSRNSTPGSSRGNSPARMASGGGETALALLLLDRLNQLESKVSGKGQ	
Qy 300	241	${\tt QQQGQTVTKKSAAE} {\tt ASKKPRQKRTATKQYNVTQAFGRRGPEQTQGNFGDQDLIRQGTDYK$	
Db 300	241	QQQGQTVTKKSAAEASKKPRQKRTATKQYNVTQAFGRRGPEQTQGNFGDQDLIRQGTDYK	
Qy 360	301	${\tt HWPQIAQFAPSASAFFGMSRIGMEVTPSGTWLTYHGAIKLDDKDPQFKDNVILLNKHIDA}$	
Db 360	301	HWPQIAQFAPSASAFFGMSRIGMEVTPSGTWLTYHGAIKLDDKDPQFKDNVILLNKHIDA	
Qy 420	361	${\tt YKTFPPTEPKKDKKKKTDEAQPLPQRQKKQPTVTLLPAADMDDFSRQLQNSMSGASADST}$	
Db 420	361	YKTFPPTEPKKDKKKKTDEAQPLPQRQKKQPTVTLLPAADMDDFSRQLQNSMSGASADST	
QУ	421	QA 422	
Db 17.	421		

18. C). Regarding claims 9-10, without a particular identification, the antibody mobilization and labeling cited in claims are considered to be made directly or indirectly. Che et al. (A) disclose an immunoassay comprising to mobilize the anti-SARS N protein monoclonal antibody onto the N protein coated 96 well plates, wherein the antibody/antigen complex is finally detected by a HRP labeled secondary antibody and visualized by the substrate reacting to the

HRP labeled antibody/antigen complex in the same 96 well ELISA plate. Therefore, the disclosure by Che et al. (A) also meets the limitations of claims 9-10.

Therefore, the claimed subject matter is still anticipated by Che (A) for the reasons set forth above and the rejections should be maintained.

19.

Conclusion

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to BAO LI whose telephone number is (571)272-0904. The examiner can normally be reached on 6:30 am to 3:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mondesi Robert can be reached on 571-272-0956. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Bao Qun Li/ Examiner, Art Unit 1648